WEST Search History

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DATE: Sunday, March 21, 2004

Hide?	<u>Set</u> <u>Name</u>	Query	<u>Hit</u> Count
	DB=Pe	GPB,USPT,USOC,EPAB,JPAB,DWPI,TDBD; PLUR=YES; OP=ADJ	
	L5	L4 and (internal\$3 or inner or inside or anatomy or organ or heart or brain or lung or kidney or pelvis or leg or ankle or wrist)	34
	L4	L3 and (image or imaging or scan\$5)	62
\Box	L3	L2 and (scatter\$4 or backscatter\$4 or "back scatter\$4" or nonreflect\$4 or nonreflect\$4")	91
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	L1	Cassegrain	1742

END OF SEARCH HISTORY

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Search Results - Record(s) 1 through 34 of 34 returned.

1. Document ID: US 20030224801 A1

Using default format because multiple data bases are involved.

L5: Entry 1 of 34

File: PGPB

Dec 4, 2003

PGPUB-DOCUMENT-NUMBER: 20030224801

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030224801 A1

TITLE: High data rate wireless communication system

PUBLICATION-DATE: December 4, 2003

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47 Lovberg, John San Diego CA US Chedester, Richard Whately MA US Johnson, Paul Kihei ΗI US Slaughter, Louis Weston MA US

US-CL-CURRENT: <u>455/454</u>; <u>455/73</u>

Full: Title: Citation Front Review Classification Date Reference Sequences Attachments Claims KMC Draw Du

2. Document ID: US 20030080907 A1

L5: Entry 2 of 34

File: PGPB

May 1, 2003

PGPUB-DOCUMENT-NUMBER: 20030080907

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030080907 A1

TITLE: Easy set-up, low profile, vehicle mounted, satellite antenna

PUBLICATION-DATE: May 1, 2003

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Wang, James June-Ming San Marino CA US Sun, Paul K. Greenwich CT US Santora, Russell Geoffrey Pasadena CA US

US-CL-CURRENT: 343/713; 343/757

Eull: Title: Citation Front Review Classification Date Reference Sequences Attachments Claims KMC Draw. De

3. Document ID: US 20030080898 A1

L5: Entry 3 of 34

File: PGPB

May 1, 2003

PGPUB-DOCUMENT-NUMBER: 20030080898

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030080898 A1

TITLE: Easy set-up, low profile, vehicle mounted, in-motion tracking, satellite

antenna

PUBLICATION-DATE: May 1, 2003

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Wang, James June-Ming San Marino CA US Sun, Paul K. Greenwich CT US Santora, Russell Geoffrey Pasadena CA US

Mahon, John P. Thousand Oaks CA US

US-CL-CURRENT: 342/359; 342/383

Full | Title | Citation | Front | Review | Classification | Date | Reference | Sequences | Affectiments | Claims | KWC | Draw De

4. Document ID: US 20030053048 A1

L5: Entry 4 of 34 File: PGPB Mar 20, 2003

PGPUB-DOCUMENT-NUMBER: 20030053048

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030053048 A1

TITLE: Electron microscope and spectroscopy system

PUBLICATION-DATE: March 20, 2003

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Bennett, Robert Gloucestershire GB
Woolfrey, Andrew Mark Gloucestershire GB
Day, John Charles Clifford Bristol GB
Bewick, Angus Bristol GB

US-CL-CURRENT: 356/301

Full: Title | Citation | Front | Review | Classification | Date | Reference | Sequences | Attachments | Claims | KWIC | Draw, Do

5. Document ID: US 20030021044 A1

L5: Entry 5 of 34

File: PGPB

Jan 30, 2003

PGPUB-DOCUMENT-NUMBER: 20030021044

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030021044 A1

TITLE: Precision formed membrane surface for electromagnetic radiation

concentration and method of forming same

PUBLICATION-DATE: January 30, 2003

INVENTOR-INFORMATION:

NAME

CITY

STATE

COUNTRY

RULE-47

Dragovan, Mark W.

Chicago

IL

US

US-CL-CURRENT: <u>359/847</u>; <u>359/838</u>, <u>359/900</u>

Full Title Citation Front Review Classification Date Reference Sequences Attachments Claims RWC Draw De

6. Document ID: US 20030016539 A1

L5: Entry 6 of 34

File: PGPB

Jan 23, 2003

PGPUB-DOCUMENT-NUMBER: 20030016539

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030016539 A1

TITLE: High efficiency non-imaging optics

PUBLICATION-DATE: January 23, 2003

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47 Minano, Juan C. Madrid CA ES Benitez, Pablo Madrid TNES ES Gonzalez, Juan C. Madrid Falicoff, Waqidi US Solana Beach Caulfield, H. J. Cornersville US

US-CL-CURRENT: 362/347; 362/348

Full Trite: Citation Front Review Classification Date Reference Sequences Allachments Claims NWC Draw De

7. Document ID: US 20020182716 A1

L5: Entry 7 of 34

File: PGPB

Dec 5, 2002

PGPUB-DOCUMENT-NUMBER: 20020182716

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020182716 A1

TITLE: Support for chromophoric elements

PUBLICATION-DATE: December 5, 2002

INVENTOR-INFORMATION:

NAME

CITY

STATE

COUNTRY

RULE-47

Weisbuch, Claude

Paris

FR

Benisty, Henri

Palaiseau

FR

US-CL-CURRENT: 435/287.2; 313/483

Full Title Citation Front Review Classification Date Reference Sequences Attachments Claims KMC Draw Do

8. Document ID: US 20020176139 A1

L5: Entry 8 of 34

File: PGPB

Nov 28, 2002

RULE-47

PGPUB-DOCUMENT-NUMBER: 20020176139

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020176139 A1

TITLE: SONET capable millimeter wave communication system

PUBLICATION-DATE: November 28, 2002

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY
Slaughter, Louis	Weston	MA	US
Olsen, Randall	Carlsbad	CA	US
Phillips, Chester	Germantown	MD	US
Johnson, Paul	Kihei	HI	US
Lovberg, John	San Diego	CA	US
Tang, Kenneth Y.	Alpine	CA	US
Houghton, George	San Diego	CA	US
Kolinko, Vladimir	San Diego	CA	US
Mooney, Ryan	Kihei	HI	US

US-CL-CURRENT: 398/121; 370/328, 370/338

**Full: | Title: | Bitation | Front: | Review | Classification | Date | Reference | Sequences | Attachments | Claims | KMC | Draw De

9. Document ID: US 20020165002 A1

L5: Entry 9 of 34

File: PGPB

Nov 7, 2002

PGPUB-DOCUMENT-NUMBER: 20020165002

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020165002 A1

TITLE: Millimeter wave transceivers for high data rate wireless communication links

PUBLICATION-DATE: November 7, 2002

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47 Kolinko, Vladimir San Diego CA US Chedester, Richard Whately MA US Olsen, Randall B. Carlsbad CA US Lovberg, John San Diego CA US Tang, Kenneth Y. US Alpine CA

US-CL-CURRENT: 455/500; 455/73

Full Title Citation Front Review Classification Date Reference Sequences Attachments Claims RWC Draw De

10. Document ID: US 20020164958 A1

L5: Entry 10 of 34

File: PGPB Nov 7, 2002

PGPUB-DOCUMENT-NUMBER: 20020164958

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020164958 A1

TITLE: Millimeter wave and copper pair communication link

PUBLICATION-DATE: November 7, 2002

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47 Slaughter, Louis Weston MA US US Lambert, Thomas Makawao ΗI Nguyen, Huan Annanandale VA US Olsen, Randall Carlsbad CA US Lovberg, John San Diego CA US CA Tang, Kenneth Y. Alpine US Kolinko, Vladimir US

US-CL-CURRENT: 455/73; 455/8

Full: Title: Citation Front: Review Classification Date Beference:	Sequences Attachments Claims NMC Draw De

11. Document ID: US 20020164951 A1

L5: Entry 11 of 34

File: PGPB

Nov 7, 2002

Dec 16, 2003

PGPUB-DOCUMENT-NUMBER: 20020164951

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020164951 A1

TITLE: Millimeter wave and ethernet communication system

PUBLICATION-DATE: November 7, 2002

INVENTOR-INFORMATION:

ATE COUNTRY RULE-47
US

US-CL-CURRENT: 455/39; 455/25

Full	Title	Citation Front Review Classification Date Reference Sequences Attachments Claims KWC Draw De

	12	Document ID: US 6665546 B2

File: USPT

US-PAT-NO: 6665546

L5: Entry 12 of 34

DOCUMENT-IDENTIFIER: US 6665546 B2

TITLE: High speed, point-to-point, millimeter wave dated communication system

DATE-ISSUED: December 16, 2003

INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP	CODE	COUNTRY
Slaughter; Louis	Weston	MA			
Hill; Jon	Socorro	NM			
Lambert; Thomas	Makawao	HI			
Nguyen; Huan	Annandale	VA			
Olsen; Randall	Carlsbad	CA			
Lovberg; John	San Diego	CA			
Tang; Kenneth Y.	Alpine	CA			
Kolinko; Vladimir	San Diego	CA			

US-CL-CURRENT: 455/562.1; 370/310, 455/10, 455/25, 455/505, 455/67.15

Full Title Citation Front	Review Classification Date	Reference	Claims : KW	C Disw De

13. Document ID: US 6639733 B2

L5: Entry 13 of 34

File: USPT

Oct 28, 2003

US-PAT-NO: 6639733

DOCUMENT-IDENTIFIER: US 6639733 B2

TITLE: High efficiency non-imaging optics

DATE-ISSUED: October 28, 2003

INVENTOR-INFORMATION:

Gonzalez; Juan C.

NAME CITY STATE ZIP CODE COUNTRY

Minano; Juan C. Madrid ES
Benitez; Pablo Madrid ES

Falicoff; Waqidi Solana Beach CA

Madrid

Caulfield; H. J. Cornersville TN

US-CL-CURRENT: <u>359/728</u>; <u>359/718</u>, <u>359/726</u>, <u>362/327</u>

Full: Title: Citation: Front: Review: Classification Date: Reference: Citation Citat

14. Document ID: US 6567678 B1

L5: Entry 14 of 34 File: USPT

May 20, 2003

ES

US-PAT-NO: 6567678

DOCUMENT-IDENTIFIER: US 6567678 B1

TITLE: Multiplex sensor and method of use

DATE-ISSUED: May 20, 2003

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Oosta; Gary_M. Gurnee IL

Jeng; Tayy-Wen Vernon Hills IL

Lindberg; John M. Grayslake IL

McGlashen; Michael L. Grayslake IL

Pezzaniti; Joseph L. Round Lake IL

US-CL-CURRENT: 600/316; 356/364, 600/310

Full: Title: Chation: Front: Review Classification Date: Reference:

15. Document ID: US 6545645 B1

L5: Entry 15 of 34 File: USPT Apr 8, 2003

US-PAT-NO: 6545645

DOCUMENT-IDENTIFIER: US 6545645 B1

TITLE: Compact frequency selective reflective antenna

DATE-ISSUED: April 8, 2003

INVENTOR-INFORMATION:

NAME

CITY

STATE

ZIP CODE

COUNTRY

Wu; Te-Kao

Rancho Palos Verdes

CA

US-CL-CURRENT: 343/781P; 343/781CA

######################################	Full Title Cit	ation Front Review (lassification (Late	Reference	
	040404000000000000000000000000000000000				

16. Document ID: US 6502944 B1

L5: Entry 16 of 34

File: USPT

Jan 7, 2003

US-PAT-NO: 6502944

DOCUMENT-IDENTIFIER: US 6502944 B1

TITLE: Precision formed membrane surface for electromagnetic radiation

concentration and method of forming same

DATE-ISSUED: January 7, 2003

INVENTOR-INFORMATION:

NAME

CITY

STATE

ZIP CODE

COUNTRY

Dragovan; Mark W.

Chicago

IL

60615

US-CL-CURRENT: 359/847; 359/851, 359/857, 359/868

Full Title Citation Front Review Classification Date Reference: Classification Date Reference: Classification	DtancDe
17 17 Dociment ID: US 6449103 B1	

US-PAT-NO: 6449103

L5: Entry 17 of 34

DOCUMENT-IDENTIFIER: US 6449103 B1

TITLE: Solid catadioptric omnidirectional optical system having central coverage means which is associated with a camera, projector, medical instrument, or similar

article

DATE-ISSUED: September 10, 2002

INVENTOR-INFORMATION:

CITY

STATE

File: USPT

ZIP CODE

COUNTRY

Sep 10, 2002

Charles; Jeffrey R.

Pasadena

CA

91104

US-CL-CURRENT: 359/725; 359/366, 359/729, 359/859

Full Title Citation Front Review Classification Date Reference Citation Claims KMC Draw De

18. Document ID: US 6333826 B1

L5: Entry 18 of 34

File: USPT

Dec 25, 2001

US-PAT-NO: 6333826

DOCUMENT-IDENTIFIER: US 6333826 B1

TITLE: Omniramic optical system having central coverage means which is associated

with a camera, projector, or similar article

DATE-ISSUED: December 25, 2001

INVENTOR-INFORMATION:

NAME

CITY

STATE

ZIP CODE

COUNTRY

Charles; Jeffrey R.

Pasadena

CA

91104

US-CL-CURRENT: <u>359/725</u>; <u>359/728</u>, <u>359/729</u>, <u>359/731</u>

Full: Title: Citation Front Review Classification Date Reference Citation Claims KMC Draw De

19. Document ID: US 6070093 A

L5: Entry 19 of 34

File: USPT

May 30, 2000

US-PAT-NO: 6070093

DOCUMENT-IDENTIFIER: US 6070093 A

** See image for Certificate of Correction **

TITLE: Multiplex sensor and method of use

DATE-ISSUED: May 30, 2000

INVENTOR-INFORMATION:

Jeng; Tayy-Wen

Lindberg; John M.

NAME

Oosta; Gary M. Gurnee

Vernon Hills

CITY

Grayslake

IL

McGlashen; Michael L. Grayslake

IL

Pezzaniti; Joseph L. Round Lake IL

STATE

IL

TT.

ZIP CODE

US-CL-CURRENT: 600/316; 356/39, 600/310, 600/322

Full. Title: Citation Front Review Classification Date Reference 2008 2008 2008 Claims KNNC Draw De

COUNTRY

20. Document ID: US 6054947 A

L5: Entry 20 of 34

File: USPT

Apr 25, 2000

US-PAT-NO: 6054947

DOCUMENT-IDENTIFIER: US 6054947 A

TITLE: Helicopter rotorblade radar system

DATE-ISSUED: April 25, 2000

INVENTOR-INFORMATION:

NAME

CITY

STATE ZIP CODE

COUNTRY

Kosowsky; Lester H.

Stamford

CT

06905

US-CL-CURRENT: 342/191; 342/192, 342/25A, 342/25F, 342/25

Full Title Citation Front Review Classification Date Reference Communication Date Reference

21. Document ID: US 5578140 A

L5: Entry 21 of 34

File: USPT

Nov 26, 1996

US-PAT-NO: 5578140

DOCUMENT-IDENTIFIER: US 5578140 A

TITLE: Solar energy plant

DATE-ISSUED: November 26, 1996

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Yogev; Amnon Rehovot IL
Krupkin; Vladimir Rishon LeZion IL
Epstein; Michael Rishon LeZion IL

US-CL-CURRENT: <u>136/246</u>; <u>126/572</u>, <u>126/600</u>, <u>126/685</u>, <u>126/686</u>, <u>136/248</u>, <u>422/186</u>,

60/641.5

Full Title Citation Front Review Classificat	ion Date Reference	Claims 1000 Drave De
***************************************		***************************************
22. Document ID: US 5227797	A	
L5: Entry 22 of 34	File: USPT	Jul 13, 1993

US-PAT-NO: 5227797

DOCUMENT-IDENTIFIER: US 5227797 A

TITLE: Radar tomography

DATE-ISSUED: July 13, 1993

INVENTOR-INFORMATION:

NAME

CITY

STATE

ZIP CODE

ZIP CODE

COUNTRY

Murphy; Quentin M.

Bronxville

NY

10708

US-CL-CURRENT: 342/22; 600/425

Full: Title Citation Front Review Classification Date Reference Glassification Date Reference

23. Document ID: US 4843242 A

L5: Entry 23 of 34

File: USPT

Jun 27, 1989

US-PAT-NO: 4843242

DOCUMENT-IDENTIFIER: US 4843242 A

** See image for <u>Certificate of Correction</u> **

TITLE: Infrared microscope employing a projected field stop

DATE-ISSUED: June 27, 1989

INVENTOR-INFORMATION:

NAME

CITY

STATE

COUNTRY

Doyle; Walter M.

Laguna Beach

CA

US-CL-CURRENT: 250/330; 250/338.1, 250/341.8, 359/389

Full Title Citation Front Review Classification Date Reference Claims KinC Draw D.

24. Document ID: US 4797685 A

L5: Entry 24 of 34

File: USPT

Jan 10, 1989

US-PAT-NO: 4797685

DOCUMENT-IDENTIFIER: US 4797685 A

TITLE: Offset shaped antenna reflector

DATE-ISSUED: January 10, 1989

INVENTOR-INFORMATION:

NAME

CITY

STATE ZIP CODE COUNTRY

Guler; Michael G.

Melbourne

FL

Patel; Sharadchandra D.

Melbourne

FLFL

Conn; James K. Foster; Marcus L. Indialantic Palm Bay

FL

US-CL-CURRENT: 343/912; 29/600, 343/840

25. Document ID: US 4688325 A

L5: Entry 25 of 34

File: USPT

Aug 25, 1987

COUNTRY

US-PAT-NO: 4688325

DOCUMENT-IDENTIFIER: US 4688325 A

TITLE: Technique for fabricating offset, shaped antenna reflectors

DATE-ISSUED: August 25, 1987

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE

Guler; Michael G. Melbourne FL Patel; Sharadchandra D. Melbourne FL

Conn; James K. Indialantic FL Foster; Marcus L. Palm Bay FL

US-CL-CURRENT: 29/600

Full: Title: Chation Front: Review Classification Date Screence Management (Claims KMC) Draw Dr

26. Document ID: US 4655592 A

L5: Entry 26 of 34

File: USPT

Apr 7, 1987

US-PAT-NO: 4655592

DOCUMENT-ÍDENTIFIER: US 4655592 A

TITLE: Particle detection method and apparatus

DATE-ISSUED: April 7, 1987

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Allemand; Charly D. Newtonville MA

US-CL-CURRENT: 356/237.3; 219/121.74, 250/222.2, 250/559.16, 250/559.41, 356/338,

<u>359/859, 362/297</u>

Full: Title: Châtion Front: Review Classification Date Reference Claims KWC Disw De

☐ 27. Document ID: US 4595833 A

L5: Entry 27 of 34

File: USPT

Jun 17, 1986

US-PAT-NO: 4595833

DOCUMENT-IDENTIFIER: US 4595833 A

** See image for Certificate of Correction **

TITLE: Multiple <u>internal</u> reflection cell optical system for use in infrared spectrophotometry of liquid and fluidized samples

DATE-ISSUED: June 17, 1986

INVENTOR-INFORMATION:

NAME

CITY

Full Title Citation Front: Review Classification Date Reference

STATE

ZIP CODE

COUNTRY

Claims 0000 Draw U

Sting; Donald W.

New Canaan

CT

06840

US-CL-CURRENT: 250/353; 250/252.1, 250/338.1, 250/339.06, 250/339.11, 356/300,

<u>359/859</u>

☐ 28. Document ID: US 4404468 A

L5: Entry 28 of 34

File: USPT

Sep 13, 1983

US-PAT-NO: 4404468

DOCUMENT-IDENTIFIER: US 4404468 A

TITLE: Pyrodetector

DATE-ISSUED: September 13, 1983

INVENTOR-INFORMATION:

NAME

CITY

STATE ZIP CODE

COUNTRY

Kleinschmidt; Peter

Munich

DE

US-CL-CURRENT: 250/342; 250/338.3, 250/347, 250/349, 250/DIG.1

29. Document ID: US 4126866 A

L5: Entry 29 of 34

File: USPT

Full: Title: Citation Front: Review Classification Date Reference Carting Claims NMC Draw De

Nov 21, 1978

US-PAT-NO: 4126866

DOCUMENT-IDENTIFIER: US 4126866 A

TITLE: Space filter surface

DATE-ISSUED: November 21, 1978

INVENTOR-INFORMATION:

NAME

CITY

STATE

ZIP CODE

COUNTRY

Pelton; Edward L.

Columbus

OH

US-CL-CURRENT: 343/909; 343/872

Full Title Citation Front Review Classification Date Reference

30. Document ID: US 4106023 A

L5: Entry 30 of 34

File: USPT

Aug 8, 1978

US-PAT-NO: 4106023

DOCUMENT-IDENTIFIER: US 4106023 A

TITLE: Navigation aid system

DATE-ISSUED: August 8, 1978

INVENTOR-INFORMATION:

NAME

CITY

STATE

ZIP CODE

COUNTRY

Baghdady; Elie J.

Weston

MA

02193

US-CL-CURRENT: 342/405

Full Title: Chation Front Review Classification Date Reference Claims RMC Draw D

31. Document ID: US 3968362 A

L5: Entry 31 of 34

File: USPT

Jul 6, 1976

US-PAT-NO: 3968362

DOCUMENT-IDENTIFIER: US 3968362 A

TITLE: Optical system for laser doppler homodyne detection

DATE-ISSUED: July 6, 1976

INVENTOR-INFORMATION:

NAME CITY

STATE

ZIP CODE

COUNTRY

Mocker; Hans W.

Minneapolis

MN

Full Trifle: Citation Front Review Classification Date Reference Citation Claims RMC Disw. De

US-CL-CURRENT: 250/216; 356/5.09

32. Document ID: US 3852765 A

L5: Entry 32 of 34

File: USPT

Dec 3, 1974

US-PAT-NO: 3852765

DOCUMENT-IDENTIFIER: US 3852765 A

TITLE: SPHERICAL DOUBLE REFLECTOR ANTENNA

COUNTRY

DATE-ISSUED: December 3, 1974

INVENTOR-INFORMATION:

NAME

CITY Bresler; Aaron D.

Merrick

NY

Stein; Emanuel

Fair Lawn

NJ

STATE ZIP CODE

Erdmann; M. Otto

Denville

NJ

US-CL-CURRENT: 343/781R; 343/837, 343/912

Full Title: Chation Front Review Classification Date Reference Claims RWC Diaw Do

33. Document ID: US 3822098 A

L5: Entry 33 of 34

File: USPT

Jul 2, 1974

US-PAT-NO: 3822098

DOCUMENT-IDENTIFIER: US 3822098 A

** See image for Certificate of Correction **

TITLE: MULTISPECTRAL SENSOR MEANS MEASURING DEPOLARIZED RADIATION

DATE-ISSUED: July 2, 1974

INVENTOR-INFORMATION:

CITY STATE ZIP CODE COUNTRY NAME

Rudder; Charles L. Granite City MO Leader; John C. Manchester MO Owsley; David P. MO Florissant

Dalton; William A. J.

Florissant MO

US-CL-CURRENT: 356/320; 250/339.11, 250/347, 356/364, 356/407, 356/448

Full Title Citation Front Review Classification Date Reference District Classification Date Reference

34. Document ID: US 3814504 A

L5: Entry 34 of 34

File: USPT

Jun 4, 1974

US-PAT-NO: 3814504

DOCUMENT-IDENTIFIER: US 3814504 A

** See image for Certificate of Correction **

TITLE: REFLECTING LENS ARRAY

DATE-ISSUED: June 4, 1974

INVENTOR-INFORMATION:

CITY STATE ZIP CODE COUNTRY NAME

Brady; John Francis Clifton NJ Grill; William Augustus

Lake Parsippany

NJ

US-CL-CURRENT: 359/627; 359/727

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Гегт	Documents
INNER	3008923
INNERS	623
INSIDE	2397817
INSIDES	20991
ANATOMY	21604
ANATOMIES	640
ANATOMYS	0
ORGAN	101417
ORGANS	94467
HEART	162688
HEARTS	9109
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L5: Entry 19 of 34

File: USPT

May 30, 2000

DOCUMENT-IDENTIFIER: US 6070093 A

** See image for <u>Certificate of Correction</u> **
TITLE: Multiplex sensor and method of use

Abstract Text (3):

(b) scattering

Brief Summary Text (16):

"Noninvasive" (NI) glucose sensing techniques measure in-vivo glucose concentrations without collecting a blood sample. As defined herein, a "noninvasive" apparatus or method is one which can be used without removing a sample from, or without inserting any instrumentation into, the tissues. The concept involves irradiating a vascular region of the body with electromagnetic radiation and measuring the spectral information that results from one of four primary processes: reflection, absorption, scattering, or emission. The extent to which each of these processes occurs is dependent upon a variety of factors, including the wavelength and polarization state of the incident radiation and the glucose concentration in the body part. Glucose concentrations are determined from the spectral information by comparing the measured spectra to a calibration curve or by reference to a physical model of the tissue under examination. A brief description of noninvasive glucose measurements in the prior art is provided below.

Brief Summary Text (24):

U.S. Pat. Nos. 5,362,966, 5,237,178, 5,533,509, 4,655,225. The principles of operation are similar to those described for the NIR, except that the penetration depth of the MIR light is less than that for NIR. As a consequence, most measurements in this region have been performed using a backscattering geometry. As defined herein, a "backscattering geometry" describes a configuration wherein scattered radiation is collected on the same side of the sample as the entry point of the incident radiation. A "transmission geometry" describes a configuration wherein light is transmitted through the sample and collected on the opposite side of the sample as the entry point of the incident radiation.

Brief Summary Text (31):

Scattering

Brief Summary Text (32):

As defined herein "scattering" includes Rayleigh, Mie, and Raman scattering. Glucose decreases the intensity of Mie scattering by decreasing the refractive index difference between the extracellular fluid (ECF) and cell membranes. Gratton et al. (U.S. Pat. No. 5,497,769) have proposed a sensor based upon this effect; however, the signal to noise ratio for this technique is expected to be inadequate for glucose measurement.

Brief Summary Text (33):

Raman scattering

Brief Summary Text (34):

U.S. Pat. No. 5,553,616 teaches the use of Raman scattering with excitation in the

near infrared (780 nm) and an artificial neural network for measuring blood glucose. Glucose Raman bands that are distinct from protein Raman bands may be chosen, however, the sensitivity of this method limits its applicability for invivo measurements. WO 92/10131 discusses the application of stimulated Raman spectroscopy for detecting the presence of glucose.

Brief Summary Text (51):

A change in temperature can have a nonlinear effect on the infrared spectrum by altering the intensities as well as the frequencies of the dominant water absorption bands. A temperature change will also modify the refractive index of the sample which, in turn, will alter the scattering properties of the sample. The effective optical path length will change as a result of the aforementioned change in scattering properties. Thus, physiological and spectral parameters are often inseparably linked and a change in one of these variables can modulate the impact of other interfering variables. The result is a nonlinear change in the measured signal for a linear change in one of the physiological or spectral variables.

Brief Summary Text (57):

On a longer time scale, the physical properties of human skin change as a normal function of aging. These changes include decreased solubility (Schnider, S. L., and Kohn, R. R., J. Clin. Invest. 67, (1981) pp.1630-1635), decreased proteolytic digestibility (Hamlin, C. R., Luschin, J. H., and Kohn, R. R., Exp. Gerontol. 13, (1978) pp. 415-523), increased heat denaturation time (Snowden, J. M., Eyre, D. R., and Swann, D. H., Biochem. Biophys. Acta, 706, (1982) pp. 153-157), and the accumulation of yellow and fluorescent materials (LaBella, F. S., and Paul, G., J. Gerontol., 20, (1964) pp. 54-59). These changes appear to be accelerated in diabetes, and may alter the scattering properties of the skin via the formation of intermolecular crosslinks between collagen fibrils.

Brief Summary Text (72):

As will be described more fully below, the present invention measures the reflected, <u>scattered</u>, absorbed, emitted, or transmitted light as a function of multiple dimensions. As defined herein, a "dimension" is a measured quantity. It can be related to light which is reflected, <u>scattered</u>, absorbed, emitted, or transmitted by the sample. It can also be related to time or space or both.

Brief Summary Text (80):

(b) scattering

Brief Summary Text (87):

(b) scattering

Brief Summary Text (94):

(b) scattering

Brief Summary Text (101):

(b) scattering

Brief Summary Text (106):

Another embodiment of the present invention provides enhanced selectivity by illuminating the sample with electromagnetic radiation and recording the intensity of the reflected, absorbed, <u>scattered</u>, emitted or transmitted radiation as a function of at least three dimensions wherein the at least three dimensions are selected from the group consisting of:

Brief Summary Text (113):

(b) scattering

Brief Summary Text (119):

(b) scattering

Brief Summary Paragraph Table (1):

TABLE 1

Char. Frequ. (Hz).sup.a Vis. Raman.sup.b NIR Raman.sup.b Fluorescence.sup.b NIR.sup.b MIR.sup.b Polarization.sup.b Photoacoustics.sup.b

Subject

Temperature 0.1-1 6 6 5 2 2 3 2 pH 0.01 3 3 5 3 3 5 5 Tissue Scattering 10-100 1 3 5 3 3 8 8 Pulsatile flow 1 5 5 5 7 4 8 8 Body part movement 1-10 2 4 4 1 1 3 3 Electrolyte Concentrations 0.1-1 3 3 5 3 3 8 4 Pressure at the interface .1-10 6 6 3 3 3 5 3 Refractive Index 0.1-10 5 4 3 3 3 5 3

Legend .sup.a Numerals in this column indicate the oscillation frequencies (Hz) of several important physiological variables. .sup.b Numerals in this column indicate the relative sensitivity (1-10, 1 being most sensitive) of the spectroscopic measurement to several important physiological variables.

Drawing Description Text (6):

FIGS. 5(a)-5(c) are is a depiction of the <u>scattered</u> light intensity predicted for a sample which interacts with the light produced by the polarization modulator of FIG. 4, wherein the sample behaves as: 5(a) a linear polarizer oriented at 45.degree., 5(b) a linear polarizer oriented at 0.degree., and 5(c) a circular polarizer.

Detailed Description Text (4):

When electromagnetic radiation impinges on a sample, the radiation is reflected, scattered, absorbed, emitted, or transmitted. The extent to which any of these processes occurs depends upon the chemical constitution of the sample as well as the frequency (or wavelength), polarization state, and angle of incidence of the impinging light beam. The methods of the prior art employ only a small fraction of the available spectroscopic information for measurement purposes. As a result, they are unable to accurately measure glucose in the presence of dominating contributions from physiological and spectral interferences.

Detailed Description Text (14):

##EQU2## (b) Secondly, the photoacoustic response is a function of the optical energy absorbed (as opposed to transmitted). Thus, scattering effects are much less important for PAS measurements than for optical absorbance measurements.

<u>Detailed Description Text</u> (16): Scattering

Detailed Description Text (17):

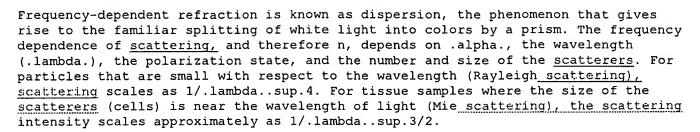
An electromagnetic wave incident on an isolated molecule with an electron cloud will cause the electrons to oscillate about their equilibrium positions, in synchrony with the applied wave. The resulting electronic oscillator emits radiation (scatters) in all directions in a plane perpendicular to the oscillating electrons. Some molecules are more susceptible to applied electromagnetic waves than others, and the tendency of their electrons to oscillate is defined by a parameter, a, called the polarizability.

Detailed Description Text (18):

Refraction is the result of radiation <u>scattered</u> in the same direction as that of the incident light wave. The phase of the <u>scattered</u> wave is different from that of the wave that passes through the sample. These two types of waves then recombine (interfere) to produce a wave that has apparently passed through the sample with a different velocity. The parameter used to describe this phenomenon is called the refractive index (n) defined as:

<u>Detailed Description Text</u> (20):

The refractive index, and therefore the scattering, is dependent on frequency.



Detailed Description Text (21):

In tissues, light <u>scattering</u> occurs because of a mismatch between the index of refraction of the ECF and the cells and organelles comprising the tissue. The index of refraction of the ECF varies as its composition changes, whereas the index of the cellular membranes and organelles remains relatively constant.

<u>Detailed Description Text</u> (22):

A change in the glucose concentration in tissues can alter the intensity and directionality of <u>scattering</u>; however, a direct measurement of glucose via scattering is difficult because:

Detailed Description Text (23):

(a) the change in <u>scattering</u> produced by a physiologically relevant change in glucose concentration is extremely weak,

Detailed Description Text (24):

(b) a change in glucose concentration can initiate a complex interplay of hormonally-regulated metabolic reactions in the intra- and extracellular fluid, the products of which reactions may also alter scattering intensity, and

Detailed Description Text (25):

(c) a change in glucose concentration may also alter the size of the cells (and thus their scattering properties) via a change in osmolarity of the ECF.

Detailed Description Text (26):

Raman Scattering

<u>Detailed Description Text</u> (27):

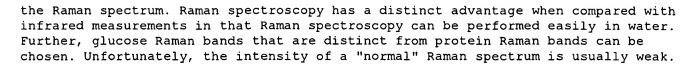
When light impinges on a sample, most of the <u>scattered</u> photons are elastically (or Rayleigh) <u>scattered</u>, meaning that they have the same frequency as the incident radiation. A small fraction of the <u>scattered</u> light (approximately one in a thousand incident photons) is inelastically <u>scattered</u> at frequencies that are shifted by frequencies defined by molecular vibrations. Raman <u>scattering</u> occurs at frequencies corresponding to the incident frequency plus or minus a molecular vibrational frequency as shown below:

Detailed Description Text (28):

where .nu..sub.Raman is the Raman <u>scattered</u> frequency, .nu..sub.o is the incident (laser) frequency and v.sub.vib is a vibrational frequency of the molecule under study. Raman bands having frequencies that are lower than the incident frequency are called "Stokes" shifted bands. Those with frequencies that are higher than the incident frequency are called "Anti-Stokes" bands. Stokes and Anti-Stokes shifted Raman bands are displaced symmetrically about the incident frequency. A Raman spectrum is thus a vibrational spectrum that is obtained by recording the intensity of scattered light as a function of frequency.

Detailed Description Text (29):

Because the selection rules for Raman <u>scattering</u> are different from those of MIR or NIR, Raman <u>scattering</u> is complementary to these other techniques. In other words, vibrational modes that produce intense Raman bands may be invisible in the MIR or NIR spectra. Additionally, some IR and NIR vibrational bands may not be present in



Detailed Description Text (30):

If the laser excitation frequency lies within an electronic absorption band of a chromophore in the molecule, then some vibrations associated within that chromophore can be dramatically enhanced. This technique, known as Resonance Raman scattering, can be used to increase the sensitivity of the measurement. For example, NIR excitation into the heme absorption bands will cause an enhancement of the heme vibrational modes, far above the intensity of the protein Raman bands. Normal and Resonance Raman spectroscopies may thus yield vibrational information that is complementary to that obtained by IR or NIR methods.

Detailed Description Text (33):

Chiral molecules, such as glucose, are molecules that cannot be superimposed on their mirror <u>image</u>. A unique property of chiral molecules is that they are optically active, i.e., they have different refractive indexes for Left- and Right-circularly polarized light. Their differential interaction with Left- and Right-circularly polarized light is measured by means of a technique called polarimetry. Polarimetry is a method of measuring and describing changes in the polarization state of light upon interaction with a polarization element (e.g. a sample). The polarization properties of a polarization element can be divided into three groups: (1) diattenuation, (2) retardance, and (3) depolarization (see, for example, J. L. Pezzaniti, Mueller Matrix <u>Imaging</u> Polarimetry, Dissertation, 1993).

Detailed Description Text (35):

There are subtle, but important differences between diattenuation and dichroism. Diattenuation is measured as the difference in scattered intensity between two orthogonal polarization states. The orthogonal polarization states might be Left-circularly polarized versus Right-circularly polarized states (for a circular diattenuator) or Vertical-polarized versus Horizontally-polarized states (for a linear diattenuator). In diattenuation, the scattered intensity is modulated by both absorption and scattering processes. This contrasts with dichroism, which measures the difference in the amount of light absorbed by two orthogonal polarization states.

Detailed Description Text (36):

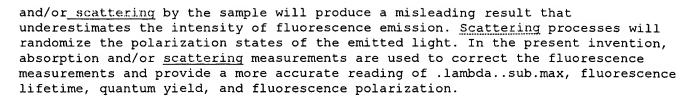
For turbid samples, e.g. biological samples, <u>scattering</u> effects are important and diattenuation measurements are performed. For weakly <u>scattering</u> samples, diattenuation typically reduces to the simpler circular dichroism measurement.

Detailed Description Text (40):

Depolarization is a process in which completely polarized light is coupled to unpolarized light and is defined as ##EQU4## In turbid media, an incident polarized light beam undergoes multiple scattering events. The polarization of the incident beam is degraded with each scattering event, and the depolarization can be used as a measure of the number of scattering events in the medium. Because glucose influences the overall refractive index in tissue, the number of scattering interactions changes with varying glucose concentrations, because scattering interactions increases, the polarized light becomes increasingly depolarized. Thus as glucose concentration changes, the scatter distribution changes and the depolarization can be monitored as an indirect measurement of scatter and glucose concentration.

Detailed Description Text (54):

All of the fluorescence parameters described above may be used to measure analyte properties (such as concentration or dynamics), however, all of these methods suffer from absorption and <u>scattering</u> interferences in the sample. Absorption



<u>Detailed Description Text</u> (57): Corrections for tissue scattering

Detailed Description Text (58):

Light <u>scattering</u> depends on the wavelength and the polarization state of the incident light as well as the difference in refractive index between the <u>scattering</u> center(s) and the surrounding medium. For tissue samples, light <u>scattering</u> arises from the refractive index mismatch between cell or mitochondrial membranes, collagen fibers or other organelles, and the exctracellular fluid (ECF) of the tissue. As used herein, tissue <u>scattering</u> is taken to mean light that is <u>scattered</u> by tissues. Tissue <u>scattering</u> can change over temporal or spatial dimensions due to a variety of factors, such as the water distribution or collagen content in the tissues, diet, or disease states, such as diabetes or hypertension. For example, an increase of water in the tissue ECF will decrease the index of refraction, thereby increasing the difference in refractive index between the ECF and the cell membranes, thereby increasing tissue <u>scattering</u>. Tissue <u>scattering</u> can vary considerably between individuals due to changes in skin properties, disease states, diet or even exercise.

Detailed Description Text (59):

Tissue <u>scattering</u> can lead to spurious or nonlinear results for spectroscopic measurements for a variety of reasons. First, <u>scattering</u> results in a loss of transmitted intensity due to <u>scattering</u> at angles that are outside the numerical aperture of the collection optics. For an absorbance measurement, the effect is a misleading reading that overestimates the amount of light absorbed. Secondly, multiple <u>scatter</u> events within the tissue lead to an ill-defined optical path length. Photon trajectories through the tissue are not rectilinear as they are in homogenous media. Multiple reflections and refractions effectively increase the optical path length.

Detailed Description Text (60):

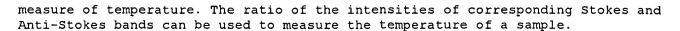
The polarization measurements described above can be used to compensate for (normalize) the effects of <u>scattering</u> on spectroscopic measurements. Diattenuation, retardance, and depolarization measurements provide a complete description of the polarization properties, and therefore the refractive index, of the sample. In particular, depolarization may be used as a measure of person-to-person or day-to-day variation in <u>scattering</u>. Multiple wavelengths may be used to enhance selectivity.

<u>Detailed Description Text</u> (61):

Absorbance measurements that are sensitive to <u>scattering</u> or optical path length can be corrected by using known relationships between the refractive index and the <u>scattering</u> properties of the medium. For example, the empirical relationship between wavelength and <u>scatter</u> intensity may be used to estimate the extent of <u>scatter</u> at wavelengths where direct measurement of <u>scatter</u> are difficult. For example, polarimetric measurements described above can be performed in the visible region or NIR region and used to normalize the measurements made at other wavelengths, e.g. measurements in MIR region.

Detailed Description Text (65):

The temperature dependence of the MIR water spectrum complicates glucose measurements by providing a variable background signal that must be subtracted in order to reveal the glucose information. Raman <u>scattering</u> can be an accurate



Detailed Description Text (66):

Spatial dimensions may also be used to sort out the contributions from pH, electrolytes, and temperature. A temperature gradient exists between the outer surface of the skin and the tissue <u>inside</u>. Accordingly, the effects of temperature on the measured signal should scale with the distance into the tissue (normal to the tissue surface) whereas contributions from electrolyte concentrations and pH should be more evenly distributed. Thus, in the foregoing example, a spatial dimension is used to separate out the effects of temperature on the measured signal from those effects due to pH and electrolyte concentrations.

Detailed Description Text (72):

For noninvasive measurements of in-vivo parameters, the incident radiation must pass through the stratum corneum before reaching viable tissues, and hence the thickness, composition, and morphology of the stratum corneum can affect measurements (see FIG. 2). As the beam penetrates into the tissue, the radiation may be scattered, absorbed, reflected or emitted by structures and chromophores that can vary dynamically and between individuals. As a result, the beam intensity decreases rapidly as it penetrates into the tissue and the majority of the spectroscopic signal arises near the surface of the tissue where the light intensity is at a maximum. As defined herein, the penetration depth, d, is the distance within the tissue at which the incident light intensity (I.sub.o) falls to (I.sub.o /e).

Detailed Description Text (73):

The <u>scattering</u>, absorption, and emission properties vary with the wavelength and the polarization state of the incident light. Light of different wavelengths may reach vastly different penetration depths within the tissue, and essentially all noninvasive measurements are dependent on wavelength and polarization. This effect can be used to advantage, since a judicious choice of wavelength and polarization state can provide a level of control over the depth to which the tissue will be probed by the radiation. This control can be used to selectively extract information related to particular chromophores, based on their predictable, or measurable, spatial locations within the sample.

<u>Detailed Description Text</u> (76):

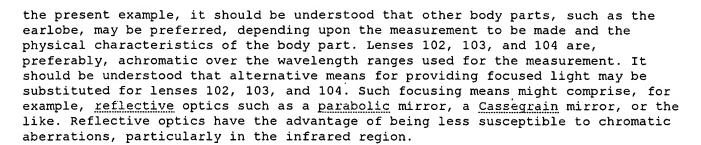
Spatial dimensions that are generally parallel to the skin surface can also be used to provide additional selectivity. Spectroscopic \underline{images} of biological tissues generally contain regular repeating structures. \underline{Image} analysis techniques (such as multidimensional Fourier transformation, segmentation, or some other \underline{image} processing technique) can be used to extract signals contained in certain locations or spatial frequencies in the \underline{image} .

Detailed Description Text (77):

For example, a spectroscopic <u>image</u> of a tissue sample may reveal blood vessels of regular sizes and separation distances. Such structures will contribute particular spatial frequencies to the <u>image</u>. Selective spectroscopic information may be obtained from these structures by collecting a spectroscopic <u>image</u> (i.e. recording a spectroscopic variable as a function of two spatial variables) and performing a multidimensional Fourier transform on the spectroscopic <u>image</u>. The signals resulting from the blood vessels would then be selectively obtained by measuring the spectroscopic signal intensity as a function of spatial frequency. An additional advantage of the foregoing method is that the measured signal is relatively insensitive to changes in alignment or sample positioning.

Detailed Description Text (82):

The filtered light is focused onto a body part, e.g. a finger 117, by a focusing means comprising, for example, lenses 103 and 104. While a finger has been used in



Detailed Description Text (83):

Lenses 102, 103, and 104 may be held in a fixed position or, alternatively, may be moved in order to alter the focal point within the tissue or at the tissue surface. Such an altered focal point might comprise, for example, a focal point that is translated along an axis normal to the skin surface or in a direction parallel to the skin surface. For example, such a moving focal point of light might be used to make measurements as a function of the spatial position of the focused light beam. Alternatively, a defocused beam of light may be used to minimize the power density at the tissue surface or for imaging applications where it is desirable to distribute light evenly across the tissue sample.

Detailed Description Text (93):

Normalization of the acoustic signal may be accomplished by a number of alternative means. For example, the measured photoacoustic signal may be normalized to the intensity of the light emitted by light source 115. Alternatively, a measurement of the scattered intensity may be made at detector 122 (as described below).

Detailed Description Text (96):

Referring again to FIG. 3, the light exiting the polarization modulator is reflected from a patterned mirror 113 and focused onto the finger. The purpose of this beam is to measure the depolarization and diattenuation of the finger (or other body part, such as an earlobe). Measurements of the intensity of the scattered light are made by the detectors 160, 161, and 162, which are placed close to or, preferably, in contact with the finger 117. A circular polarizer is placed immediately before the active area of each detector. The circular polarizer transmits right-circular polarized light and blocks left-circular polarized light. As defined herein, close to the skin means within about 0 to 10 mm from the surface of the skin, preferably from about 0 to 1 mm from the surface of the skin or, most preferably, in contact with the skin. The placement of detectors 160, 161, and 162 close to or, preferably, in contact with the skin provides several advantages including:

Detailed Description Text (97):

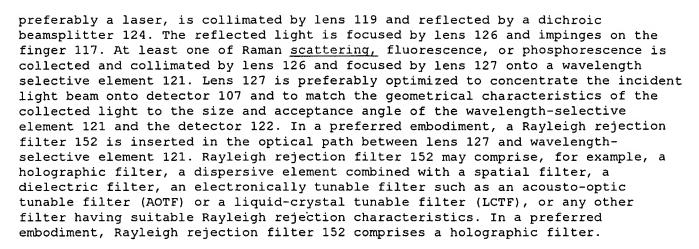
(1) the capability of resolving small angular and spatial distributions of the scattered light,

<u>Detailed Description Text</u> (100):

As shown in FIG. 3, three detectors 160, 161, and 162 are used to detect the backscattered radiation; however, it is to be understood that a different number of detectors can be used. Detectors 160, 161, and 162 generate electrical signals that are representative of the intensity and polarization state of the scattered radiation. These electrical signals can be analyzed by means of a lock-in amplifier that is tuned to the modulation frequency of the polarization modulator or harmonics of that modulation frequency. Alternatively, the electrical signals produced by the detectors 160, 161, and 162 can be digitized and analyzed by means of a digital filter, such as a Fourier Transform.

Detailed Description Text (104):

Returning again to FIG. 3, the present invention provides a means for performing Raman and emission measurements as follows. Light from a light source 115,



<u>Detailed Description Text</u> (105):

Wavelength-selective element 121 allows certain wavelengths of light to be transmitted to the detector 122 by means of either a dispersive or interferometric selection mechanism. Wavelength selective elements, such as a Czerny-Turner monochromator, can be used in a scanning mode with a point detector or, preferably, the wavelength selective element is coupled to an array detector. A typical array detector may be a silicon photodiode array or, in a preferred embodiment, the array detector may be a charge coupled device (CCD) or Charge Injection Device (CID) detector. InGaAs detectors are optimized for the NIR and can be operated at room temperature or cooled to liquid nitrogen temperatures.

Detailed Description Text (115):

As shown in FIG. 6, a single detector 114 measures the intensity of the scattered radiation as a function of the incident polarization state. Detector 114 may comprise a point detector or an imaging detector, such as a CCD, a CID. In a preferred embodiment, detector 114 is a point detector. A point detector, such as a photodiode, has a faster response time than the CCD and, therefore, is more amenable to high frequency modulation and detection schemes. In an alternative embodiment, the scattered light is measured using an imaging detector, such as a CCD or a CID. An imaging detector has the capability of recording the scattered intensity as a function of one or more spatial dimensions along the skin surface.

Detailed Description Text (117):

In a preferred embodiment, polarization state analyzer 123 is a simple thin film polarizer, such as a stretched polymeric film, positioned in front of the detector 108. The polarizer may be either a linear polarizer or a circular polarizer. In a particularly preferred configuration, depolarization is measured by modulating the incident polarization state as shown in FIG. 4, and analyzing the scattered light with a fixed polarizer.

Detailed Description Text (118):

Although a finger is shown in FIG. 6, it should be understood that other body parts, such as an earlobe, may be preferably used for depolarization measurements in order to increase the sensitivity of the measurement. For example, the <u>internal</u> structures (bones, cartilage, tendons) of the finger will almost completely depolarize the light which is incident on the finger. The polarization state of light passing through the earlobe, however, is preserved to a greater degree due to the lower amount of <u>internal</u> structure in the earlobe relative to the finger.

Detailed Description Text (123):

The apparatus of FIG. 8 comprises a multispectral <u>imaging</u> system that uses multiple spectral dimensions (i.e. IR, Raman, Fluorescence, etc.) and one or more spatial dimensions. Additionally, a series of measurements may be performed over time, thereby adding a temporal dimension.

<u>Detailed Description Text</u> (125):

image field and is achromatic.

Detailed Description Text (126):

The system shown in FIG. 8 exhibits several advantage. First, multiple spectroscopic <u>images</u> may be recorded. The system shown in FIG. 8 might, for example, be used to measure a temporal oscillation in one or more parameters of the sample.

Detailed Description Text (127):

By using <u>imaging</u> detectors and <u>image</u> analysis techniques, such as a multidimensional Fourier transform, the signals that are contained in particular spatial frequencies across an <u>image</u> plane of the tissue sample may be selected. For example, a spectroscopic <u>image</u> of a tissue sample will contain blood vessels of regular sizes and such structures will contribute particular spatial frequencies to the <u>image</u>. Spectroscopic information may be obtained from these structures by collecting a spectroscopic <u>image</u> (i.e., an <u>image</u> at multiple wavelengths) and performing a multidimensional Fourier transform on the spectroscopic <u>image</u>. Other image processing techniques, such as segmentation, may also be employed.

<u>Detailed Description Text</u> (130):

An exploded view of another embodiment of the present invention is provided in FIG. 9. Device 201 is a hand-held, noninvasive multiplex sensor that may be used for measuring at least one parameter (e.g. the concentration of an analyte such as glucose), in a sample, (e.g. a body part). An activating button 202 is depressed by the user to activate the instrument prior to use. Optical head 203 contains an array of light sources and detectors and an optical window 204, which is transparent to the light that is emitted or detected by device 201. Appropriate light sources may be, for example, infrared emitting diodes (IREDs) or laser diodes. Photodetectors 207, which are also provided inside the optical head, measure the light that is <u>backscattered</u> by the sample. Preferred photodetectors may be, for example, a photodiode, a charge coupled device, or a charge injection device. Any other suitable detector may be used. The detectors are preferably optimized for a given wavelength range and may consist of, for example, silicon, InGaAs, Ge, or PbS detectors. Photodetectors 207 are attached to a preamplifier board 208, which contains electronic circuitry in accordance with its function. Batteries 206 provide power for the device. Optical window 204 could also be designed to filter out stray ambient light, thereby further reducing noise in the measurement.

Detailed Description Text (131):

Several preselected wavelengths of infrared radiation are focused onto the sample. Radiation that is reflected, emitted or <u>scattered</u> by the sample is collected by detectors 207. Quantitative analysis is performed by the central processing unit 215 in conjunction with a multivariate calibration model and algorithms stored in module 217. A concentration value is subsequently displayed by a display unit 212, which is connected to the central processing unit 215. Display unit 212 is preferably a liquid crystal display, which is large enough to be easily read by patients with visual dysfunction, such as that caused by advanced stage diabetes. Alternatively, an audible readout may be provided.

Detailed Description Text (133):

One embodiment of the present invention provides a method of measuring in-vivo glucose concentrations that combines measurements of infrared absorbance (in both forward and <u>backscattering</u> geometries) with diattenuation measurements. The results of these measurements are shown in Table 3 below.

Detailed Description Text (136):

Another embodiment of the present invention provides a method of measuring blood

glucose that is corrected for spectral variables (such as water (tissue hydration), hemoglobin, tissue <u>scattering</u> (refractive index), and temperature) by means of a combination of complementary spectroscopic techniques. Specifically, glucose measurements are performed using a combination of infrared absorbance, photoacoustics, and scattering measurements.

Detailed Description Text (139):

A sample is illuminated with infrared light and the infrared absorbance is measured as a function of the wavelength of light absorbed. A plot of the infrared absorbance versus wavelength is referred to as an infrared absorbance spectrum and has an associated spectral dimension, namely, the wavelength of the light absorbed by the sample. Within the same embodiment, at least one additional complementary spectroscopic technique is applied, (e.g. Raman scattering, photoacoustics, polarimetry, fluorescence spectroscopy, etc.), thereby adding at least one more spectral dimension to the measurement.

Detailed Description Text (140):

Further, and within the same embodiment, at least two spectral dimensions are recorded as a function of at least one spatial dimension of the sample. In the present example, a spatial dimension of the sample may be measured using an imaging detector, such as a charge-coupled device (CCD) detector, or by multiple discrete detectors. Further, and still within the same embodiment, the measurements described above are performed over time in order to measure a temporal dimension. The concentration of the at least two sample constituents are thus measured as a function of at least two spectral dimensions, at least one spatial dimension, and at least one temporal dimension.

<u>Detailed Description Paragraph Table</u> (1): TABLE 3	Wavelength Measurement (nm) % CV
1. IR,	Forward Scattering 1000 nm 26.1 2.
IR, Forward Scattering 1150 nm 25.9 3. IR, Ba	ackscattering 1400 nm 23.8 4. IR,
Backscattering 1650 nm 23.1 5. DC scatter 633	3 nm 26.2 6. Circular diattenuation 633
nm 25.6 7. 1-4 combined 17.2 8. 1-6 combined	16.5

CLAIMS:

- 1. A method for measuring at least one parameter of a sample comprising the steps of:
- (a) illuminating said sample with light;
- (b) performing at least two spectroscopic measurements, wherein said at least two spectroscopic measurements are different members of the group consisting of:

infrared absorbance,

scattering,

diattenuation,

emission spectroscopy,

photoacoustic spectroscopy,

provided that said photoacoustic spectroscopy relates acoustic signal intensity directly to the measurement of said at least one parameter of said sample; and

(c) analyzing said spectroscopic measurements to determine a measurement of said at least one parameter of said sample.

- 10. The method of claim 9, wherein said physiological variable is selected from the group consisting of:
- (i) temperature,
- (ii) pulsatile flow,
- (iii) tissue scattering, and
- (iv) sample heterogeneity.
- 12. A method for measuring at least one parameter of a sample comprising the steps of:
- (a) illuminating said sample with light;
- (b) performing at least two spectroscopic measurements, wherein said at least two spectroscopic measurements are different members of the group consisting of:

infrared absorbance,

scattering,

polarimetry,

emission spectroscopy,

photoacoustic spectroscopy,

provided that said photoacoustic spectroscopy relates acoustic signal intensity directly to the measurement of said at least one parameter of said sample; and

- (c) analyzing said spectroscopic measurements to determine a measurement of said at least one parameter of said sample, further comprising the step of measuring at least one spatial dimension.
- 13. A method for measuring at least one parameter of a sample comprising the steps of:
- (a) illuminating said sample with light;
- (b) performing at least two spectroscopic measurements, wherein said at least two spectroscopic measurements are different members of the group consisting of:

infrared absorbance,

scattering,

polarimetry,

emission spectroscopy,

photoacoustic spectroscopy,

provided that said photoacoustic spectroscopy relates acoustic signal intensity directly to the measurement of said at least one parameter of said sample; and

(c) analyzing said spectroscopic measurements to determine a measurement of said at least one parameter of said sample, further comprising the step of measuring at

least one temporal dimension.

- 14. A method for measuring at least one parameter of a sample comprising the steps of:
- (a) illuminating said sample with light;
- (b) performing at least two spectroscopic measurements, wherein said at least two spectroscopic measurements are different members of the group consisting of:

infrared absorbance,

scattering,

polarimetry,

emission spectroscopy,

photoacoustic spectroscopy,

provided that said photoacoustic spectroscopy relates acoustic signal intensity directly to the measurement of said at least one parameter of said sample; and

- (c) analyzing said spectroscopic measurements to determine a measurement of said at least one parameter of said sample, further comprising the step of measuring at least one spatial dimension and at least one temporal dimension.
- 15. A method for measuring at least one parameter of a sample comprising the steps of:
- (a) illuminating said sample with light;
- (b) performing at least two spectroscopic measurements, wherein said at least two spectroscopic measurements are different members of the group consisting of:

infrared absorbance,

scattering,

polarimetry,

emission spectroscopy,

photoacoustic spectroscopy,

provided that said photoacoustic spectroscopy relates acoustic signal intensity directly to the measurement of said at least one parameter of said sample; and

(c) analyzing said spectroscopic measurements to determine a measurement of said at least one parameter of said sample, further comprising the step of removing an interference related to at least one spectral variable.